

## CLAIMS

1. A transgenic non-human mammal or a portion thereof, wherein an  $\alpha$ -synuclein gene is introduced and the gene is expressed in the neurons, and the number of dopamine-producing neurons in the substantia nigra is significantly decreased as compared with that of a wild-type animal.
2. The transgenic non-human mammal or a portion thereof according to claim 1, wherein the  $\alpha$ -synuclein gene is a human  $\alpha$ -synuclein gene or a variant thereof.
3. The transgenic non-human mammal or a portion thereof according to claim 1 or 2, wherein the  $\alpha$ -synuclein gene is a variant of a wild-type human  $\alpha$ -synuclein gene in a manner that substitutes a Thr residue for an Ala residue at amino acid residue 53 in an amino acid sequence encoded by the wild-type human  $\alpha$ -synuclein gene.
4. The transgenic non-human mammal or a portion thereof according to any of claims 1 to 3, wherein the  $\alpha$ -synuclein gene is a gene that is varied from a wild-type  $\alpha$ -synuclein gene in a manner that deletes C terminal amino acid residues encoded by the wild-type  $\alpha$ -synuclein gene.
5. The transgenic non-human mammal or a portion thereof according to any of claims 1 to 4, wherein a recombinant DNA incorporating the  $\alpha$ -synuclein gene therein under the control of a promoter capable of expressing the  $\alpha$ -synuclein gene in the dopamine-producing neurons is introduced.
6. The transgenic non-human mammal or a portion thereof according to any of claims 1 to 5, wherein the promoter capable of expressing the  $\alpha$ -synuclein gene in the dopamine-producing neurons is a tyrosine hydroxylase promoter.
7. The transgenic non-human mammal or a portion thereof according to any of claims 1 to 6, wherein an intracerebral dopamine level at an early age is significantly decreased as compared with that of a wild-type animal.
8. The transgenic non-human mammal or a portion thereof according to any of claims 1 to 7, wherein an intracerebral dopamine level at an early age is decreased to 85% or less as compared with that of a wild-type animal.

9. The transgenic non-human mammal or a portion thereof according to any of claims 1 to 8, wherein a tyrosine hydroxylase expression level is decreased to 80% or less as compared with that of a wild-type animal.

10. The transgenic non-human mammal or a portion thereof according to any of claims 1 to 9, wherein a spontaneous locomotor activity is decreased to 60% or less as compared with that of a wild-type animal.

11. The transgenic non-human mammal or a portion thereof according to any of claims 1 to 10, wherein the non-human mammal is a mouse.

12. A method for screening a substance having dopamine-like action wherein the non-human mammal or a portion thereof according to any of claims 1 to 11 is used.

13. The screening method according to claim 12, wherein the substance having dopamine-like action is a therapeutic agent or preventive agent for Parkinson's disease.

14. A substance obtained by the screening method according to claim 12 or 13.

15. A therapeutic agent or preventive agent for Parkinson's disease which comprises a substance obtained by the screening method according to claim 12 or 13, as an active ingredient.